1/2 ページ

Nucleotide

"Exhibit A"

Display Settings: GenBank

Gm1 promoter and use thereof

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GenBank: DD249890.1
FASTA Graphics
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VERSION
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            WO 2005108574-A/1.
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            Sciurognathi; Muroidea; Muridae; Murinae; Mus; Mus.
REFERENCE
            1 (bases 1 to 3871)
  AUTHORS
          Oeda, K. and Takahashi, Y.
  TTTLE
            Gml promoter and use thereof
  JOURNAL
           Patent: WO 2005108574-A 1 17-NOV-2005;
            SUMITOMO CHEMICAL COMPANY LIMITED
COMMENT
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                WO 2005108574-A/1
            PD
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            PF
                 15-MAR-2005 WO 2005JP005077
            PR
                15-MAR-2004 JP 04P 072244
            PT
                kenji oeda, yasuhiko takahashi
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2/2 ページ

Gm1 promoter and use thereof - Nucleotide result

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3841 gcctgcttgg ggcgctctca ggtgcgagct c



IPI.

GenCore version 6.3 Copyright (c) 1993 - 2009 Biocceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on: September 16, 2009, 05:20:38; Search time 3986 Seconds

(without alignments)

98157.397 Million cell updates/sec

Title: US-10-593-216-1

Perfect score: 3871

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Gapop 10.0 , Gapext 1.0

Searched: 87455243 seqs, 50535937365 residues

Total number of hits satisfying chosen parameters: 174910486

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Scoring table: IDENTITY NUC

Post-processing: Minimum Match 0%

Maximum Match 100% Listing first 45 summaries

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SUMMARIES

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REFERENCE
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 AUTHORS
           Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
            Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
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           Mouse whole genome scaffolding with paired end reads from 10kb
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  JOURNAL
           Unpublished (2000)
COMMENT
           Contact: Robert B. Weiss
           University of Utah Genome Center
            University of Utah
           Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
           84112, USA
           Tel: 801 585 5606
            Fax: 801 585 7177
```

Email: ddunn@genetics.utah.edu

FEATURES

ORTGIN

Db

Db

Οv

3/21 ページ

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(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptored DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gi|4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and

purified. The sheared, adaptored mouse DNA was annealed to adaptored vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

Query Match 16.8%; Score 649; DB 20; Length 649; Best Local Similarity 100.0%;

Matches 649; Conservative 0: Mismatches 0: Indels O: Gaps

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1169 CGAAGGTCCTGAGTTCAAATACCAGCAACCACATGGTGGCTCACAACCATCTGTAATGGG 1228

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          Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
           Sciurognathi; Muroidea; Muridae; Murinae; Mus.
REFERENCE
           1 (bases 1 to 589)
 AUTHORS
           Dunn, D., Aovagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
          Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
          Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von
           Niederhausern, A. and Wright, D., Weiss, R.
 TITLE
          Mouse whole genome scaffolding with paired end reads from 10kb
           plasmid inserts
 JOURNAL
           Unpublished (2000)
COMMENT
           Contact: Robert B. Weiss
           University of Utah Genome Center
           University of Utah
           Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC. UT
          84112, USA
          Tel: 801 585 5606
          Fax: 801 585 7177
          Email: ddunn@genetics.utah.edu
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                   (http://www.jax.org/resources/documents/dnares/). The DNA
                  was hydrodynamically sheared by repeated passage through a
                  0.005 inch orifice at constant velocity. The sheared DNA
                  was blunt end-repaired with T4 DNA polymerase and T4
                  polynucleotide kinase. Adaptor oligonucleotides were
                  ligated to the blunt ends in high molar excess. The
                  adaptored DNA was purified and size-selected for a 9.5 to
                  10.5 kb range using preparative agarose gel
                  electrophoresis. Vector DNA was prepared from a derivative
                  of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
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